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Stimulation of the Noradrenergic System during Memory Formation Impairs Extinction Learning but not the Disruption of Reconsolidation

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The noradrenergic system plays a critical role in the 'consolidation' of emotional memory. If we are to target 'reconsolidation' in patients with anxiety disorders, the noradrenergic strengthening of fear memory should *not* impair the disruption of reconsolidation. In Experiment I, we addressed this issue using a differential fear conditioning procedure allowing selective reactivation of one of two fear associations. First, we strengthened fear memory by administering an α_2-adrenergic receptor antagonist (ie, yohimbine HCl; double-blind placebo-controlled study) 30 min before acquisition (time for peak value yohimbine HCl < 1 h). Next, the reconsolidation of one of the fear associations was manipulated by administering a β-adrenergic receptor antagonist (ie, propranolol HCl) 90 min before its selective reactivation (time for peak value propranolol HCl < 2 h). In Experiment II, we administered propranolol HCl *after* reactivation of the memory to rule out a possible effect of the pharmacological manipulation on the memory retrieval itself. The excessive release of *noradrenaline* during memory formation not only delayed the process of extinction 48 h later, but also triggered broader fear generalization. Yet, the β-adrenergic receptor blocker during reconsolidation *selectively* 'neutralized' the fear-arousing aspects of the noradrenergic-strengthened memory and undermined the generalization of fear. We observed a similar reduction in fear responding when propranolol HCl was administered *after* reactivation of the memory. The present findings demonstrate the involvement of noradrenergic modulation in the formation as well as generalization of *human* fear memory. Given that the noradrenergic strengthening of fear memory impaired extinction learning but not the disruption of reconsolidation, our findings may have implications for the treatment of anxiety disorders.
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Keywords: fear memory; reconsolidation; extinction; generalization; noradrenergic modulation; anxiety disorders

INTRODUCTION

The process of reconsolidation, the protein-synthesis-dependent restabilization of a memory upon retrieval, enables the modification of memory representation (Nader *et al*, 2000). We previously demonstrated that disrupting reconsolidation by administering the β-adrenergic receptor antagonist propranolol HCl before reactivation *selectively* 'deleted' the emotional expression of a fear memory in humans (ie, startle fear responding) (Kindt *et al*, 2009; Soeter and Kindt, 2010, 2011a). Importantly, the fear-erasing effects following the pharmacological manipulation were not restricted to the feared cue itself, but instead generalized to category-related information (Soeter and

Kindt, 2011a). Given that *fear generalization* lies at the heart of many anxiety disorders (Lissek *et al*, 2008), targeting the process of reconsolidation may provide a novel therapeutic strategy in the treatment of (for instance) post-traumatic stress disorder. However, there are a number of conditions that may prevent reconsolidation from occurring, such as the strength of memories (Suzuki *et al*, 2004; Wang *et al*, 2009). If we are to target reconsolidation in patients suffering from post-traumatic stress disorder, strong fear memory should evidently not act as a constraint on reconsolidation.

To date, a substantial body of evidence supports the noradrenergic modulation in the formation of emotional memory (McGaugh and Roozendaal, 2009). Recently, we demonstrated that stimulation of the noradrenergic system during memory formation strengthened the emotional expression of *human* associative fear memory (Soeter and Kindt, 2011b). That is, the excessive release of noradrenaline during memory formation not only delayed the 'process of extinction' 48 h later, but also generated a superior recovery of fear following *re*-exposure to the original

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stressor. These findings thus suggest that noradrenaline may play an important role in the etiology and maintenance of anxiety disorders. However, the effect of noradrenaline on fear generalization (ie, a main characteristic of anxiety disorders) currently remains unknown. The generalization of fear seems to be dependent on the strength of the memory as operationalized by training intensity (Laxmi et al, 2003). Hence, an interesting question is whether the noradrenaline-induced strengthening of associative fear memory also promotes the generalization of fear responding. At the same time, the stimulation of the noradrenergic system during memory formation should *not* impair the disruption of reconsolidation if we are to target reconsolidation clinically.

Here, we addressed these issues by using a differential fear conditioning procedure allowing selective reactivation of one of two categorically distinct fear associations sharing the same aversive outcome and a test of fear generalization. Testing included different phases across three consecutive days, each separated by 24 h (see Figure 1a). During acquisition (day 1), two fear-relevant stimuli ($CS1_{Spider}$–$CS2_{Gun}$) were repeatedly paired with an aversive electric stimulus (US), whereas a fear-irrelevant stimulus was not ($CS3_{Mug}$) (Supplementary Figure S1). Furthermore—in Experiment I—the formation of fear memory was manipulated by the systemic administration of yohimbine HCl (day 1), an α_2-adrenergic receptor antagonist supposed to stimulate central noradrenergic activity by blocking the α_2-adrenergic autoreceptor (Charney et al, 1987; Peskind et al, 1995; Soeter and Kindt, 2011b). To reach peak plasma levels upon completion of the acquisition phase, participants ($n = 30$) received, in a double-blind manner, an oral dose of either 20 mg of yohimbine HCl or placebo pill 30 min before fear learning (day 1) (time for peak value yohimbine HCl <1 h; Grasing et al, 1996). On day 2, the reconsolidation of one of the fear associations (CS1) was manipulated by the systemic administration of propranolol HCl—a β-adrenergic receptor antagonist that indirectly targets the protein synthesis required for reconsolidation by inhibiting the noradrenaline-stimulated cAMP response element binding (CREB)

phosphorylation (Thonberg et al, 2002). In view of the peak plasma concentrations of propranolol HCl (Gilman and Goodman, 1996), all of the participants received, in a single-blind manner, an oral dose of 40 mg of propranolol HCl 90 min before the selective reactivation of the CS1 memory (day 2). (In line with animal studies (eg, Debiec and LeDoux, 2004), we previously demonstrated that (1) a reactivation trial in combination with placebo pill and (2) the omission of memory reactivation after propranolol HCl intake yielded intact fear responding (Kindt et al, 2009; Soeter and Kindt, 2010). Taken together, these findings indicate that both the oral administration of propranolol HCl and the reactivation of the fear memory are necessary for the observed fear-erasing effects.). Administering pills before reactivation does not—however—rule out a possible effect of the pharmacological manipulation on the retrieval of the fear memory itself. Therefore, the participants in Experiment II ($n = 10$) received, in a single-blind manner, an oral dose of propranolol HCl subsequent to memory reactivation. Memory retention was tested 24 h later (ie, first test trial day 3), followed by an *extinction* procedure, *reminder shock*, and a test of fear generalization. Expression of fear was measured using *startle fear potentiation* (Hamm and Weike, 2005). We obtained *skin conductance responding* and *retrospective US expectancy ratings* to assess the anticipation of threat (Weike et al, 2007). Here, the US expectancy was measured retrospectively instead of 'online' (Soeter and Kindt, 2010, 2011a, b) to prevent the expectancy ratings from interfering with the measurement of electrodermal activity. Salivary α-amylase and blood pressure levels were determined to ensure that the drug manipulations exerted its intended physiological effect (Stegeren et al, 2006, 2009). See Supplementary Materials and methods for a detailed description of the apparatus and materials. We hypothesized that the yohimbine HCl manipulation during acquisition would strengthen the fear memory and thereby (1) act as a constraint on the reconsolidation of the CS1 fear association, (2) delay the process of extinction learning, and (3) promote the generalization of fear responding, relative to placebo pill.
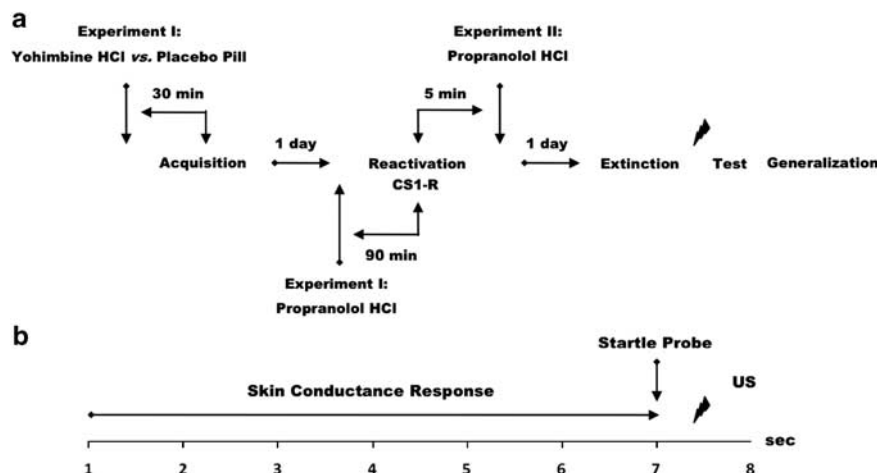

Figure 1 Schematic of the experimental design (a) and the CS1$^+$ or CS2$^+$ conditioning trial (b). In the CS1$^-$, CS2$^-$, CS3$^-$, and CS1-R trials, no US was delivered. CS, conditioned stimuli.

MATERIALS AND METHODS

Participants

In all, 40 undergraduate students (10 men, 30 women) from the University of Amsterdam ranging in the age of 18–26 years (mean ± SD age, 20.7 ± 1.9 years) participated in the study. Participants were assessed to be free from any current or previous medical or psychiatric condition that would contraindicate taking a single dose of yohimbine HCl (20 mg) and propranolol HCl (40 mg) (ie, pregnancy; seizure disorder; respiratory disorder; cardiovascular disease; blood pressure $\leq 90/60$ or $\geq 140/90$ mmHg; diabetes; liver/kidney disorder; depression; or psychosis). To eliminate individuals who might have difficulty with any temporary symptoms induced by either drug manipulation, an additional exclusion criterion contained a score ≥ 26 on the Anxiety Sensitivity Index (ASI) (Peterson and Reiss, 1992). All participants who were enrolled completed the study. The participants in Experiment I were randomly assigned to one of two conditions, with the restriction that conditions were matched on Trait Anxiety (STAI-T) (Spielberger *et al*, 1970), Spider Phobic Questionnaire (SPQ) (Klorman *et al*, 1974), and ASI scores as close as possible (see Table 1). Participants received either partial course credits or were paid a small amount (€42, –) for their participation in one of the experiments. The ethical committee of the University of Amsterdam approved the study and informed consent was obtained from all participants.

Experimental Procedure

Participants were subjected to a differential fear conditioning procedure, with three pictures serving as conditioned stimuli (CS1$_{Spider}$–CS2$_{Gun}$–CS3$_{Mug}$). The two fear-relevant pictures (CS1, CS2) were paired with an electric stimulus, whereas the fear-irrelevant picture (CS3) was not. We employed fear-relevant stimuli because they lead to a superior conditioning of aversive associations and are especially resistant to extinction learning compared with fear-irrelevant cues (Mineka and Öhman, 2002; Lang *et al*, 2005). Moreover, given that most anxiety disorders are associated with these categories of stimuli (Mineka and Öhman, 2002), we are specifically interested in targeting stronger fear memory. One of the fear-relevant stimuli (CS2) served as control for the other fear-relevant stimulus (CS1). However, since fear-relevant stimuli are known to

have an innate prepotency to elicit fear responses (Lovibond *et al*, 1994), we employed an additional fear-irrelevant control stimulus (CS3) to verify whether the procedure was capable of neutralizing the acquired fear responding. Testing included several phases across three subsequent days, each separated by 24 h. During each session, participants sat behind a table with a computer monitor at a distance of 50 cm in a sound-attenuated room. Each session began with a 1-min acclimation period consisting of 70 dB broadband noise, which continued throughout the session as background noise, followed by a habituation phase consisting of 10 startle probes to reduce initial startle reactivity. Characteristics of the CSs, trial order, ITIs, and startle probes during memory reactivation (day 2) and extinction, testing (day 3) were similar to acquisition (day 1). Assignment of the pictures as CS1$^+$ and CS2$^+$ was counterbalanced across participants.

Acquisition—day 1. Before participants subscribed to the study, they were already informed about the pill administrations and the electric stimulus. Upon arrival at the first testing day, details of the various study procedures (eg, EMG electrodes, electric stimulus, side effects of the pharmacological treatments, medical screening) were explained and possible questions were answered. Participants who agreed to enroll in the study were interviewed with regard to their health and any medical or psychiatric conditions that would contraindicate taking a single dose of yohimbine HCl (20 mg) and propranolol HCl (40 mg). In addition, blood pressure was measured. Once a participant was medically cleared, written informed consent was obtained and the ASI, SPQ, and STAI were administered. Furthermore, saliva samples were collected. To this end, participants were instructed just to place the swab in their mouths for 3 min.

After attachment of the startle, skin conductance, and shock electrodes, the intensity of the US was determined. Starting at an intensity of 1 mA, the level of a 2-ms aversive electric stimulus delivered to the wrist of the non-preferred hand was gradually increased. The intensity of shock was individually set at a level defined by the participant as 'uncomfortable, but not painful' and remained set to this intensity throughout the following days. To test for the potential effects of yohimbine HCl on the fear-potentiated startle reflex (Davis *et al*, 1993), 10 baseline startle probes (noise alone; NA) were presented before pill intake. Afterwards, participants were detached from the experimental setup and received, in a double-blind manner, an oral dose of either 20 mg of yohimbine HCl or placebo pill. To reach peak plasma levels upon completion of the acquisition phase (Grasing *et al*, 1996), a resting period of 30 min was inserted. Participants were offered magazines to read. Before acquisition, participants were attached to the experimental apparatus and were informed with regard to the CSs. They were instructed that two of the pictures would be followed by an electric stimulus in most of the cases, whereas the third picture would never be followed by the US. They were told to learn to predict whether an electric stimulus would occur or not on the basis of the three pictures.

In the acquisition phase, the CS1$^+$, CS2$^+$, and CS3$^-$ were presented five times for 8 s. The startle probe was presented

Table I Mean Values (SD) of the Reported Spider Fear, Trait Anxiety, Anxiety Sensitivity, Shock Intensity, and US Evaluation for the Yohimbine HCl and Placebo Pill Condition and the Propranolol after Reactivation Group

	Yohimbine HCl propranolol	Placebo pill propranolol	Propranolol after reactivation	T-test
Spider fear	4.6 (3.3)	6.3 (5.3)	4.2 (4.5)	$ts_{28} < 1.04$
Trait anxiety	37.5 (8.0)	36.3 (7.9)	32.0 (10.1)	$ts_{28} < 1.53$
Anxiety sensitivity	9.2 (5.4)	8.5 (5.0)	7.8 (5.6)	$ts_{28} < 1$
Shock intensity	16.4 (6.9)	14.5 (4.7)	17.3 (5.6)	$ts_{28} < 1.35$
US evaluation	−2.7 (1.0)	−2.8 (0.6)	−2.4 (0.9)	$ts_{28} < 1$

7 s after CS onset and was followed by the US 500 ms later (CS1$^+$ and CS2$^+$) (see Figure 1b). To reduce the possibility that the reactivation trial on day 2 resulted in extinction learning, the first presentation of both the CS1$^+$ and CS2$^+$ was unreinforced (LaBar et al, 1998). Five baseline startle probes were presented alone (NA). Order of trial type was randomized within blocks (ie, CS1$^+$, CS2$^+$, CS3$^-$, and NA). Intertrial intervals (ITI) varied between 15, 20, and 25 s, with a mean of 20 s.

The STAI-S was filled out both before and upon completion of the acquisition phase. In addition, blood pressure as well as saliva samples were collected. At the conclusion of the experiment, participants were asked to evaluate the pleasantness of the US. Furthermore, they were explicitly instructed to remember what they had learned during acquisition. These instructions were included to enhance retention of the CS–US contingency on the following days (Norrholm et al, 2006) and to prevent participants from erroneously expecting a different contingency scheme during subsequent testing. The procedure in Experiment II, *Propranolol after Reactivation*, was similar to Experiment I, except for pill administration and resting period.

Memory reactivation—day 2. To substantiate consolidation of the fear memory, a break of 24 h after acquisition was inserted. All of the participants received, in a single-blind manner, an oral dose of 40 mg of propranolol HCl 90 min before memory reactivation (CS1-R)—even though they were informed that they could also receive a placebo pill. The STAI-S was filled out both before pill administration and upon completion of the experiment. In addition, at these time points, blood pressure and saliva samples were collected.

After electrode attachment, participants were told that the same three pictures would be presented and they were asked to remember what they had learned during acquisition. They were again instructed that two of the pictures would be followed by an electric stimulus in most of the cases, whereas the third picture would never be followed by the US. In the memory reactivation phase, a single unreinforced CS1-R was presented for 8 s, followed by a startle probe presented alone. The procedure in Experiment II, Propranolol after Reactivation, was similar to Experiment I, except for pill administration, which occurred 5 min *after* reactivation of the memory.

Extinction, testing—day 3. Upon arriving at the experimental site, blood pressure and saliva samples were again collected. In addition, the STAI-S was completed. After attachment of the electrodes, the participants were informed that the same three pictures provided during acquisition would be presented. No further instructions were given. In the extinction phase, participants were exposed to the three pictures (CS1$^-$, CS2$^-$, and CS3$^-$) for 10 times without the electric stimulus (US). Startle probes were again presented 7 s after CS onset. Furthermore, 10 startle probes were presented alone (NA). After the extinction procedure, we presented an unsignaled reminder shock to reinstate the expression of the original fear memory. Evidence for a reinstatement effect is indicated by an increase of the differential conditioned response from the last extinction trial to the first trial at test. We predicted that the unsignaled reminder shock would not result in a return of

fear to the reactivated stimulus (CS1) given that propranolol HCl is supposed to disrupt the reconsolidation of the reactivated fear memory (CS1–US). We further predicted that yohimbine HCl would enhance the return of fear (CS2) following the unsignaled reminder shock (US) (see also, Soeter and Kindt, 2011b). Therefore, we presented only one as opposed to the traditional procedure of *three* unsignaled USs to avoid a ceiling effect in the return of fear for the participants who received a placebo pill during fear acquisition (day 1) (Norrholm et al, 2006; Kindt et al, 2009; Soeter and Kindt, 2010). The time between the last extinction trial and the reinstating US was 19 s. Following the unsignaled US, participants were again presented with 1 CS1$^-$, CS2$^-$, CS3$^-$, and NA trial (ie, reinstatement testing). The time between the reinstating USs and reinstatement testing was 18 s. Next, generalization testing took place. That is, participants were exposed to category-related *pictures* (G$_p$CS1, G$_p$CS2, and G$_p$CS3) and category-related *words* (G$_w$CS1, G$_w$CS2, and G$_w$CS3) (see Supplementary Figure S1). The order of generalization stimuli (ie, pictures *vs* words) was counterbalanced across participants. At the conclusion of the experiment, participants completed the STAI-S and judged the pleasantness of the US. In addition, participants were asked to indicate for each phase (beginning *vs* end) of the experiment to what extent they had expected the US after each of the CSs. The procedure in Experiment II, Propranolol after Reactivation, was similar to Experiment I.

Statistical Analysis

State anxiety, salivary α-amylase, and systolic as well as diastolic blood pressure were subjected to a 2 (condition: yohimbine HCl *vs* placebo pill) $\times$ 2 (moment: before *vs* after pill intake) mixed analysis of variance (ANOVA). Startle responses, electrodermal activity, and US expectancy ratings were analyzed by means of a mixed ANOVA for repeated measures with condition (ie, yohimbine HCl *vs* placebo pill) as between-subjects factor and stimulus (ie, simple contrasts: CS1 *vs* CS3 and CS2 *vs* CS3) and trial (ie, stimulus presentation) as within-subjects factors. The differential response (CS1 *vs* CS3 and CS2 *vs* CS3) was compared over testing phases, respectively (first trial *vs* last trial). To determine the *speed* of extinction learning, a 2 (condition: yohimbine HCl *vs* placebo pill) $\times$ 2 (stimulus: CS1 *vs* CS3 and CS2 *vs* CS3) $\times$ 5 (trials: averaging over each two consecutive extinction trials) mixed ANOVA was performed. To test whether the two conditioned stimuli (ie, spider and gun) were equally affected by the drug manipulation, we performed a 2 (trial: stimulus presentation) $\times$ 2 (stimulus: CS1 *vs* CS3) $\times$ 2 (stimulus category: spider *vs* gun) mixed ANOVA. Planned comparisons between the CS1 and CS2 stimuli were performed separately. Missing data due to artifacts (ie, 1.13% of the trials) were not replaced and hence excluded from the analyses. Significance was set at $p < 0.05$.

RESULTS

Manipulation Check Drug Administration

Analysis of the effect of the yohimbine HCl manipulation on blood pressure and sAA level during fear acquisition (day 1)

revealed the expected increase in both systolic and diastolic blood pressure (moment × condition, $F_{1,28} = 26.58$, $p < 0.001$, $\eta_p^2 = 0.49$; $F_{1,28} = 12.24$, $p < 0.01$, $\eta_p^2 = 0.30$, respectively), as well as salivary α-amylase (Stegeren et al, 2009) (moment × condition, $F_{1,23} = 6.50$, $p < 0.05$, $\eta_p^2 = 0.22$) in comparison to placebo pill. However, the yohimbine HCl manipulation during fear acquisition did not affect the reported state anxiety that was assessed before and upon completion of the acquisition phase (moment × condition, $F_{1,28} < 1.45$) (see also, Table 2).

The propranolol HCl manipulation during memory reactivation (day 2) did not differentially affect the BP and sAA between conditions (moment × condition, $F_{1,28} < 1.92$). In both the yohimbine and placebo pill group, we observed a significant decrease in systolic and diastolic BP (moment, $F_{1,28} = 92.22$, $p < 0.001$, $\eta_p^2 = 0.77$; $F_{1,28} = 39.81$, $p < 0.001$, $\eta_p^2 = 0.59$, respectively), as well as in the amylase level (Stegeren et al, 2006) (moment, $F_{1,22} = 7.95$, $p = 0.01$, $\eta_p^2 = 0.27$) following propranolol HCl administration, indicating that the pill manipulation exerted its intended physiological effect. Consistent with other studies (Grillon et al, 2004), the propranolol HCl manipulation during memory retrieval did not affect the reported state anxiety that was assessed before and upon completion of the reactivation phase (moment, $F_{1,28} < 1$; moment × condition, $F_{1,28} < 1$). The decrease in BP, sAA, and reported state anxiety during memory reactivation (day 2) in Experiment II, Propranolol after Reactivation, did not differ from both the yohimbine HCl and placebo pill condition (moment × condition, $F_{1,28} < 1.98$) (see also, Table 3).

Experiment I, Yohimbine HCl vs Placebo Pill

Startle fear responding. Acquisition—day 1 ANOVA showed fear conditioning on day 1 by a significant increase of the differential startle response (ie, simple contrasts: $CS1_{Spider}$ vs $CS3_{Mug}$–$CS2_{Gun}$ vs $CS3_{Mug}$) from trials 1 to 5 (stimulus × trial, $F_{1,28} = 21.73$, $p < 0.001$, $\eta_p^2 = 0.44$; $F_{1,28} = 19.85$, $p < 0.001$, $\eta_p^2 = 0.42$, respectively). We observed no difference in responses to the first trial of acquisition ($CS1_{Spider}$ vs $CS2_{Gun}$ vs $CS3_{Mug}$) (stimulus, $F_{2,27} < 1$), indicating that the fear relevancy of the pictures did not affect startle fear responding in the absence of associative

Table 2 Mean Values (SD) of the Systolic and Diastolic Blood Pressure (in mmHg) and Amylase Level (in U/ml) Pre- and Post-Pill Intake during Fear Acquisition for the Yohimbine HCl and Placebo Pill Condition (ie, Experiment I)

Fear acquisition yohimbine HCl	Pre-pill intake	Post-pill intake	T-test: two-tailed
Yohimbine condition			
Systolic BP	126.1 (SD = 10.1)	138.0 (SD = 11.9)	$t_{14} = -4.84$, $p < 0.001$
Diastolic BP	72.5 (SD = 6.3)	77.7 (SD = 8.3)	$t_{14} = -2.72$, $p < 0.05$
sAA Level	91.4 (SD = 127.1)	168.9 (SD = 233.9)	$t_{11} = -2.36$, $p < 0.05$
Placebo pill condition			
Systolic BP	125.3 (SD = 14.7)	120.8 (SD = 8.9)	$t_{14} = 2.22$, $p < 0.05$
Diastolic BP	73.0 (SD = 7.5)	70.3 (SD = 7.0)	$t_{14} = 2.25$, $p < 0.05$
sAA Level	103.1 (SD = 105.2)	75.7 (SD = 62.9)	$t_{12} < 1$

Table 3 Mean Values (SD) of the Systolic and Diastolic Blood Pressure (in mmHg) and Amylase Level (in U/ml) Pre- and Post Propranolol HCl Administration during Memory Reactivation for the Yohimbine HCl and Placebo Pill Condition (ie, Experiment I) and the Propranolol after Reactivation Group (ie, Experiment II)

Memory reactivation propranolol HCl	Pre-pill intake	Post-pill intake	T-test: two-tailed
Yohimbine condition			
Systolic BP	130.3 (SD = 11.2)	113.3 (SD = 11.7)	$t_{14} = 6.52$, $p < 0.001$
Diastolic BP	75.0 (SD = 7.6)	67.8 (SD = 6.7)	$t_{14} = 4.56$, $p < 0.001$
sAA Level	193.0 (SD = 261.4)	32.1 (SD = 38.4)	$t_{11} = 2.21$, $p < 0.05$
Placebo pill condition			
Systolic BP	125.5 (SD = 9.2)	110.3 (SD = 7.8)	$t_{14} = 7.53$, $p < 0.001$
Diastolic BP	70.1 (SD = 5.6)	65.8 (SD = 5.5)	$t_{14} = 4.72$, $p < 0.05$
sAA Level	96.1 (SD = 105.6)	41.2 (SD = 38.0)	$t_{11} = 2.32$, $p < 0.05$
Propranolol after reactivation			
Systolic BP	121.3 (SD = 4.9)	107.1 (SD = 7.0)	$t_9 = 5.94$, $p < 0.001$
Diastolic BP	73.2 (SD = 8.7)	67.8 (SD = 8.1)	$t_9 = 3.28$, $p < 0.05$
sAA Level	56.9 (SD = 41.8)	30.1 (SD = 36.2)	$t_9 = 2.55$, $p < 0.05$

learning (see also, Soeter and Kindt, 2011a). Moreover, fear responses to the reinforced pictures ($CS1_{Spider}$ vs $CS2_{Gun}$) were equally acquired (stimulus × trial, $F_{1,28} < 1$; Figure 2a and b). Furthermore, the administration of yohimbine HCl did not directly affect the fear learning given that we observed no difference during acquisition between the two conditions (stimulus × trial × condition, $F_{1,28} < 2.09$). Yohimbine HCl did also not affect the startle response *per se* as we found no effect on the habituation trials that were presented before and after pill administration (moment × trial × condition, $F_{9,20} < 1.06$).

Memory reactivation—day 2 The two groups (yohimbine HCl vs placebo pill) expressed similar levels of differential startle potentiation (CS1-R vs NA) during memory reactivation (day 2) (stimulus × condition, $F_{1,28} < 1$). Since the NA trial always followed the reactivation trial (CS1-R) on day 2, we further compared the CS1-R with the last trial of habituation (ie, an NA trial). This analysis also revealed a similar level of startle potentiation during memory reactivation for both conditions (stimulus × condition, $F_{1,28} < 1.46$). Furthermore, the absence of a significant change in startle fear responding (CS1 vs NA) from the last trial of acquisition to memory reactivation (stimulus × trial × condition, $F_{1,28} < 1$) demonstrates that the acquired fear memory was equally well consolidated in the two groups.

Memory retention—day 3 In both the yohimbine and placebo pill group, the administration of propranolol HCl significantly decreased startle potentiation to the reactivated CS1 from the last trial of acquisition to the first extinction trial 48 h later (CS1 vs CS3; stimulus × trial, $F_{1,28} = 15.98$, $p < 0.001$, $\eta_p^2 = 0.36$), irrespective of the reactivated stimulus (ie, spider vs gun) (stimulus × trial × category, $F_{1,26} < 1.16$). We even no longer observed differential startle responding to the reactivated CS1 on the first trial of extinction learning (CS1 vs CS3; $t_{14} < 1.25$; Figure 2a and b), indicating that the yohimbine HCl
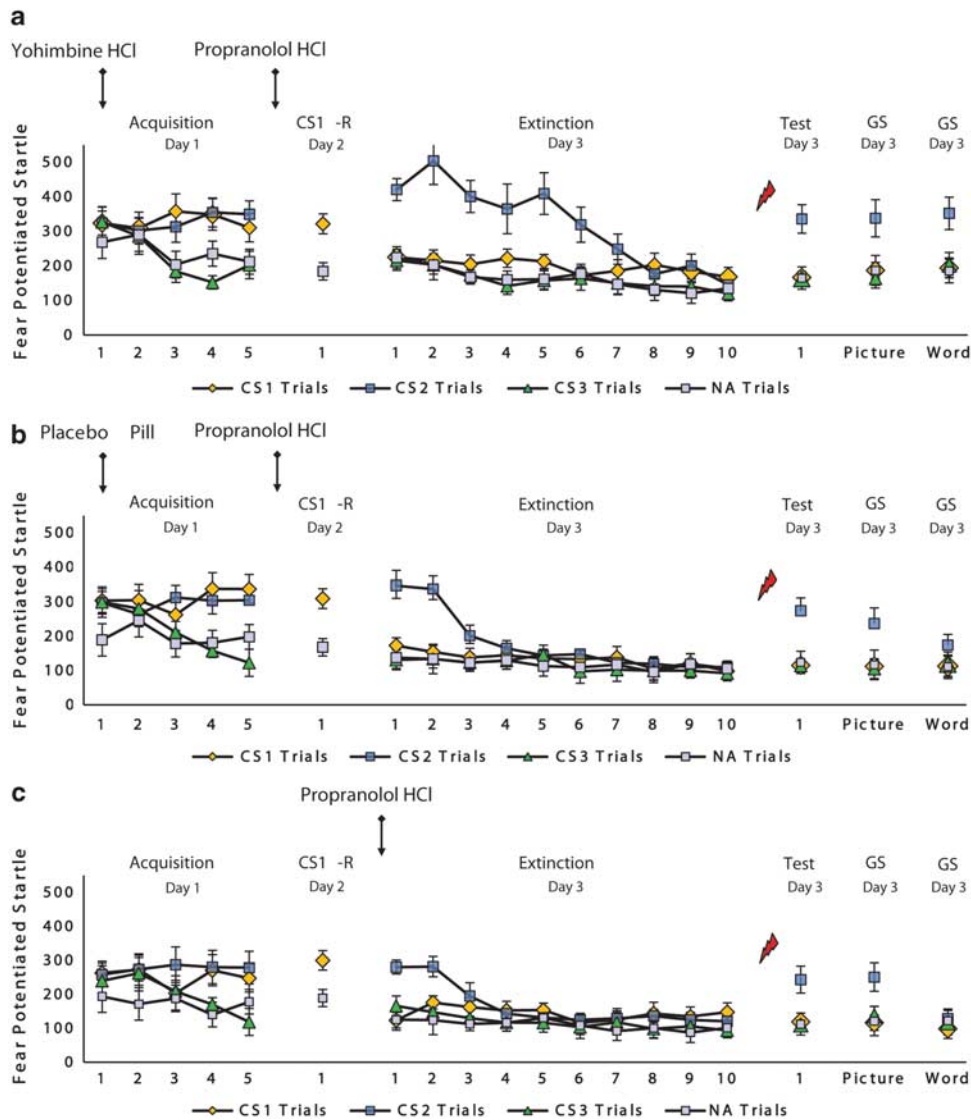
Figure 2 Mean startle potentiation to the fear-conditioned stimuli (CS1 and CS2), the control stimulus (CS3), and noise alone (NA) trials during acquisition, memory reactivation, extinction, test, and generalization for the (a) yohimbine HCl–propranolol HCl and (b) placebo pill–propranolol HCl condition (ie, Experiment I), and the (c) propranolol HCl after reactivation group (ie, Experiment II). Error bars represent SEM.

manipulation did not prevent the disruption of reconsolidation 24 h later. Conversely, startle responses to the non-reactivated CS2 remained stable from acquisition to extinction learning 48 h later in the yohimbine as well as placebo pill condition (CS2 vs CS3; stimulus × trial, $F_{1,28} < 1.12$).

Extinction learning—day 3 Given that responses to the reactivated CS1 were already eliminated on the first trial of fear extinction (day 3), we observed no extinction learning to the CS1 stimulus (CS1 vs CS3) in both the yohimbine and placebo pill group (trial 1 vs trial 10; stimulus × trial, $F_{1,28} < 1$). In contrast, startle responses to the non-reactivated CS2 decreased similarly from the first extinction trial to the last trial of extinction learning in the yohimbine and placebo pill condition (CS2 vs CS3; stimulus × trial, $F_{1,28} = 53.52$, $p < 0.001$, $\eta_P^2 = 0.66$; stimulus × trial × condition, $F_{1,28} < 1$). However, we observed a significant difference in the *speed* of the extinction learning process between the two groups (CS2 vs CS3 and CS1 vs CS2; stimulus × trial × condition, $F_{1,28} = 21.49$, $p < 0.001$,

$\eta_P^2 = 0.43$; $F_{1,28} = 23,41$, $p < 0.001$, $\eta_P^2 = 0.46$, respectively), irrespective of the non-reactivated stimulus (ie, spider vs gun) (stimulus × trial × condition × category, $F_{1,26} < 1$). Bonferroni-adjusted pairwise comparisons indeed showed that the differential startle response to the non-reactivated stimulus (CS2 vs CS3) only reached significance on the first trials of extinction learning (trials 1 and 2) in the placebo pill condition ($p < 0.001$; Figure 2b). Conversely, the differential startle response to the non-reactivated stimulus (CS2 vs CS3) remained significant up to trials 5 and 6 of extinction learning in the yohimbine group (all p's < 0.001; Figure 2a), indicating that the α_2-adrenergic drug strongly delayed the extinction learning process (see also, Soeter and Kindt, 2011b).

Reinstatement testing—day 3 Contrary to our expectations, the administration of yohimbine HCl relative to placebo pill did not result in a superior recovery of fear to the non-reactivated CS2 following the reminder shock (CS2 vs CS3; stimulus × trial × condition, $F_{1,28} < 1$). However, in both the yohimbine and placebo pill condition, the

1210

differential startle response to the non-reactivated CS2 significantly increased from the last trial of extinction to the first trial at test (stimulus × trial, $F_{1,28} = 26.56$, $p < 0.001$, $\eta_p^2 = 0.49$; Figure 2a and b). Most importantly, the reminder shock following extinction learning did not unveil a recovery of fear to the reactivated CS1 in the yohimbine as well as placebo pill condition (CS1 vs CS3; stimulus × trial, $F_{1,28} < 1.24$; CS1 vs CS3; stimulus × trial, $F_{1,28} = 36.88$, $p < 0.001$, $\eta_p^2 = 0.57$). This finding indicates that propranolol HCl effectively disrupted the reconsolidation of the reactivated fear memory and that the administration of yohimbine HCl before fear learning did not prevent this process 24 h later.

Generalization testing—day 3   We observed a significant generalization of fear to the category-related picture of the non-reactivated CS2 in both the yohimbine and placebo pill group (G_pCS2 vs G_pCS3 and G_pCS1 vs G_pCS2; main effect of stimulus, $F_{1,28} = 31.29$, $p < 0.001$, $\eta_p^2 = 0.53$; main effect of stimulus, $F_{1,28} = 25.55$, $p < 0.001$, $\eta_p^2 = 0.48$, respectively). Conversely, the category-related picture of the reactivated CS1 did not reveal any fear response in the yohimbine and placebo pill condition (G_pCS1 vs G_pCS3; main effect of stimulus, $F_{1,28} < 1.16$; Figure 2a and b). Notably, a significant difference in fear generalization for the *word* cue of the non-reactivated CS2 was observed between the two groups (G^2CS2 vs G^2CS3 and G^2CS1 vs G^2CS2; stimulus × condition, $F_{1,28} = 4.51$, $p < 0.05$, $\eta_p^2 = 0.14$; stimulus × condition, $F_{1,28} = 7.21$, $p < 0.05$, $\eta_p^2 = 0.21$, respectively). That is, we observed a significant generalization of the startle fear response to the *word* cue of the non-reactivated CS2 in the yohimbine condition (G^2CS2 vs G^2CS3 and G^2CS1 vs G^2CS2; stimulus, $F_{1,14} = 22.59$, $p < 0.001$, $\eta_p^2 = 0.62$; stimulus, $F_{1,14} = 46.86$, $p < 0.001$, $\eta_p^2 = 0.77$, respectively) but not in the placebo pill group (stimulus, $F_{1,14} < 3.19$). Hence, the yohimbine HCl manipulation triggered fear generalization to the more abstract *word* stimulus. The word cue of the reactivated CS1 did not uncover any fear response in both the yohimbine and placebo pill condition (G^2CS1 vs G^2CS3; stimulus, $F_{1,28} < 1$). Taken together, these findings demonstrate that the administration of yohimbine HCl during acquisition resulted in a broader generalization of fear 48 h later, while this generalization effect was eliminated by the administration of propranolol HCl during memory reactivation.

Analysis of the startle response to NA showed no significant differences between the two conditions during acquisition and extinction learning (condition, $F_{1,28} < 2.85$) nor during memory reactivation, reinstatement testing, and generalization ($t_{28} < 1.47$).

Skin conductance responding. Overall analysis of electrodermal responding revealed no fear conditioning during acquisition ($CS1_{Spider}$ vs $CS3_{Mug}$–$CS2_{Gun}$ vs $CS3_{Mug}$) (stimulus × trial, $F_{1,28} < 1$). When fear responses are not successfully acquired, one cannot assess the return of fear. Therefore, only subjects showing successful levels of fear acquisition (ie, trial 5 CS1 or CS2 > CS3) were included in the analyses. *Five* subjects were eliminated. That is, two subjects from the yohimbine condition and three subjects from the placebo pill group.

Subsequent analyses of variance showed no effects of the yohimbine HCl and propranolol HCl manipulation on skin conductance responding. We observed a significant increase in electrodermal activity during acquisition (trial 1 vs trial 5—day 1) in both conditions (CS1 vs CS3, CS2 vs CS3; stimulus × trial, $F_{1,23} = 11.51$, $p < 0.01$, $\eta_p^2 = 0.33$; $F_{1,23} = 7.53$, $p < 0.05$, $\eta_p^2 = 0.25$, respectively; stimulus × trial × condition, $F_{1,23} < 1$; Figure 3a and b). Furthermore, the two groups showed similar levels of electrodermal responding during the reactivation trial on day 2 (CS1-R; $t_{23} < 1$). Moreover, in both the yohimbine and placebo pill group, the differential skin conductance response (CS1 vs CS3, CS2 vs CS3) obtained during acquisition (ie, trial 5—day 1) remained stable 48 h later (ie, trial 1—day 3) (stimulus × trial, stimulus × trial × condition, $F_{1,23} < 1.52$). In both groups, a significant decrease in electrodermal responding was observed during extinction learning (trial 1 vs trial 10—day 3) (CS1 vs CS3, CS2 vs CS3; stimulus × trial, $F_{1,23} = 11.83$, $p < 0.01$, $\eta_p^2 = 0.34$; $F_{1,23} = 11.03$, $p < 0.01$, $\eta_p^2 = 0.32$, respectively; stimulus × trial × condition, $F_{1,23} < 1$). Analysis of the reinstatement effect further revealed a recovery in electrodermal responding from the last trial of extinction to the first trial at test in the yohimbine as well as placebo pill condition (CS1 vs CS3, CS2 vs CS3; stimulus × trial, $F_{1,23} = 4.77$, $p < 0.05$, $\eta_p^2 = 0.17$; $F_{1,23} = 4.35$, $p < 0.05$, $\eta_p^2 = 0.16$, respectively; stimulus × trial × condition, $F_{1,23} < 1$). In addition, the two groups showed a generalization of the skin conductance response to the category-related picture (G_pCS1 vs G_pCS3, G_pCS2 vs G_pCS3; main effect of stimulus, $F_{1,23} = 5.31$, $p < 0.05$, $\eta_p^2 = 0.19$; $F_{1,23} = 4.81$, $p < 0.05$, $\eta_p^2 = 0.17$, respectively; stimulus × condition, $F_{1,23} < 1$) as well as word cue (G_wCS1 vs G_wCS3, G_wCS2 vs G_wCS3; main effect of stimulus, $F_{1,23} = 5.08$, $p < 0.05$, $\eta_p^2 = 0.17$; $F_{1,23} = 5.56$, $p < 0.05$, $\eta_p^2 = 0.20$, respectively; stimulus × condition, $F_{1,23} < 1$). Note that analyses over the entire sample also revealed no differences between the yohimbine and placebo pill condition (CS1 vs CS3, CS2 vs CS3; stimulus × trial × condition, $F_{1,28} < 1.07$).

Retrospective US expectancy ratings. ANOVA revealed no effects of either drug manipulations on the retrospective US expectancy ratings. In both the yohimbine and placebo pill group, we observed a significant differential increase in expectancy ratings during acquisition (trial 1 vs trial 5; day 1) (CS1 vs CS3, CS2 vs CS3; stimulus × trial, $F_{1,28} = 405.18$, $p < 0.001$, $\eta_p^2 = 0.94$; $F_{1,28} = 547.74$, $p < 0.001$, $\eta_p^2 = 0.95$, respectively; stimulus × trial × condition, $F_{1,28} < 1.23$; see Supplementary Figure S2a and b). Furthermore, the two groups did not differ in their US expectancy during memory reactivation (day 2) (CS1-R; $t_{28} < 1$). Moreover, in both the yohimbine and placebo pill condition, the expectancy ratings remained stable from the last acquisition trial (day 1) to the first extinction trial 48 h later (day 3) (stimulus × trial, stimulus × trial × condition, $F_{1,28} < 1.84$). ANOVA showed extinction learning on day 3 by a significant differential decrease in US expectancy from the first extinction trial to the last trial of extinction learning in both conditions (CS1 vs CS3, CS2 vs CS3; stimulus × trial, $F_{1,28} = 208.45$, $p < 0.001$, $\eta_p^2 = 0.88$; $F_{1,28} = 165.67$, $p < 0.001$, $\eta_p^2 = 0.86$, respectively; stimulus × trial × condition, $F_{1,28} < 1.56$). We further observed a significant reinstatement effect from the last trial of extinction to the first trial at test in the yohimbine as well as placebo pill group (CS1 vs CS3, CS2 vs CS3, stimulus × trial, $F_{1,28} = 107.67$, $p < 0.001$, $\eta_p^2 = 0.79$; $F_{1,28} = 89.90$, $p < 0.001$, $\eta_p^2 = 0.76$, respectively;
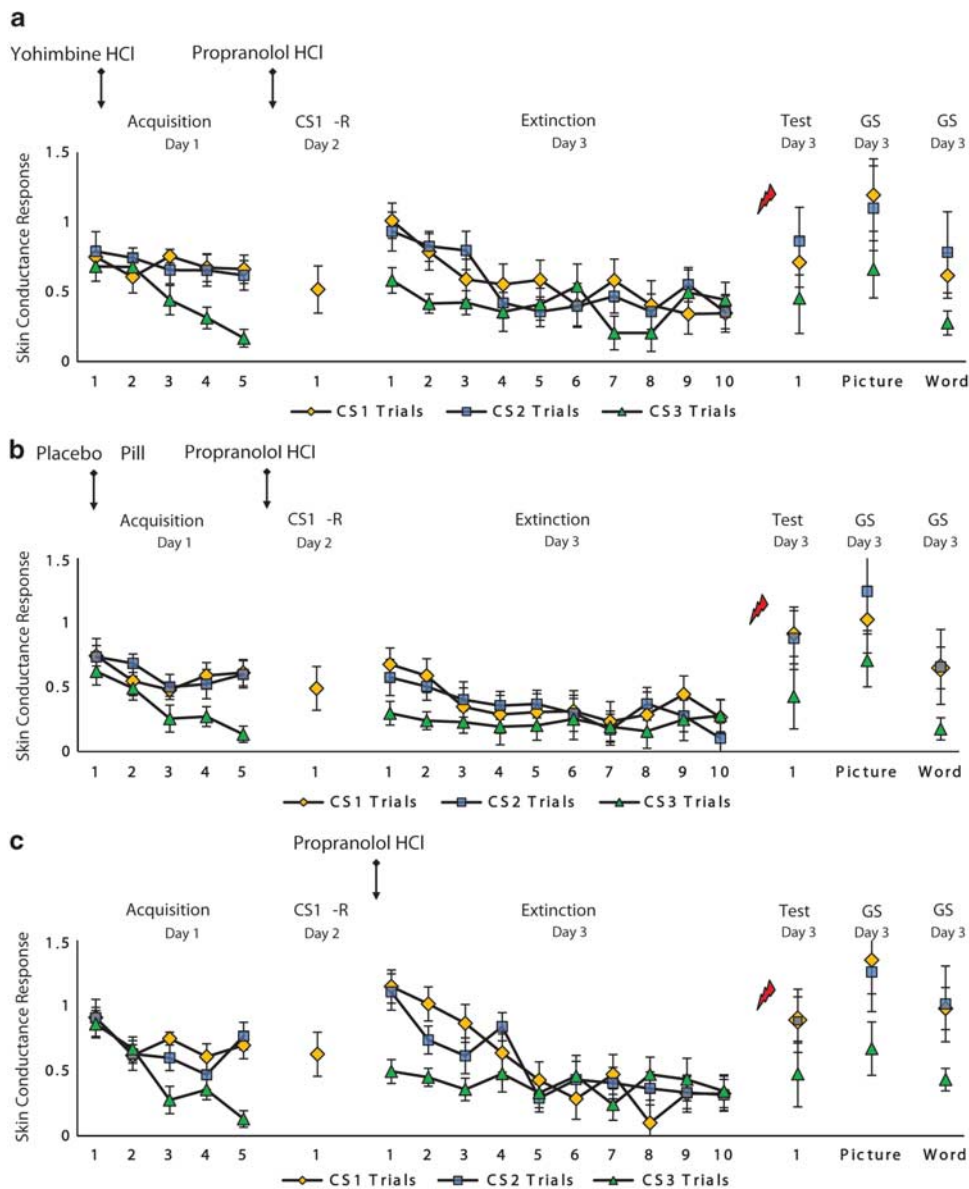
Figure 3 Mean skin conductance responses to the CS1, CS2, and CS3 trials during acquisition, memory reactivation, extinction, test, and generalization for the (a) yohimbine HCl–propranolol HCl and (b) placebo pill–propranolol HCl condition (ie, Experiment I), and the (c) propranolol HCl after reactivation group (ie, Experiment II). Error bars represent SEM.

stimulus × trial × condition, $F_{1,28} < 1.21$). Moreover, in both groups, we observed a generalization of the US expectancy ratings to the category-related picture (G_pCS1 vs G_pCS3, G_pCS2 vs G_pCS3; main effect of stimulus, $F_{1,28} = 142.73$, $p < 0.001$, $\eta_p^2 = 0.84$; $F_{1,28} = 137.87$, $p < 0.001$, $\eta_p^2 = 0.83$, respectively; stimulus × condition, $F_{1,28} < 1.45$) as well as word cue (G_wCS1 vs G_wCS3, G_wCS2 vs G_wCS3; main effect of stimulus, $F_{1,28} = 138.07$, $p < 0.001$, $\eta_p^2 = 0.83$; $F_{1,28} = 140.63$, $p < 0.001$, $\eta_p^2 = 0.83$, respectively; stimulus × condition, $F_{1,28} < 1.12$).

Experiment II, Propranolol after Reactivation

To rule out the effect of the propranolol HCl manipulation on the retrieval of the fear memory itself, participants in Experiment II received an oral dose of propranolol HCl

directly after reactivation. Resembling our previous findings (Kindt *et al*, 2009; Soeter and Kindt, 2010, 2011a), the administration of pills following memory reactivation also selectively eliminated the startle fear response 48 h after fear acquisition (CS1 vs CS3; stimulus × trial, $F_{1,9} = 12.11$, $p < 0.01$, $\eta_p^2 = 0.57$; Figure 2c), without affecting skin conductance discrimination and the retrospective US expectancy ratings (Figure 3c; see also Supplementary Results and Supplementary Figure S2c).

DISCUSSION

The present findings demonstrate that stimulation of the noradrenergic system during memory formation delayed the process of fear extinction without impairing the

disruption of reconsolidation (ie, startle fear responding). The competition between the original excitatory fear association and the newly formed inhibitory memory trace determines the behavioral outcome of extinction learning (Myers and Davis, 2002). Given that yohimbine HCl was administered during fear acquisition—and not during fear extinction 48 h later—the α_2-adrenergic drug apparently delayed the process of extinction by strengthening the original excitatory fear association (Soeter and Kindt, 2011b). Yet, the yohimbine HCl manipulation did not directly augment the differential startle fear responding either 24 or 48 h after fear learning. That is, in both the placebo pill and yohimbine HCl group, the fear responding obtained during acquisition (day 1) remained stable during memory reactivation (day 2—reactivated fear association) as well as retention testing 48 h later (non-reactivated fear association). Our finding that yohimbine HCl strengthened the original fear association without directly augmenting its behavioral expression may suggest a ceiling effect in startle fear conditioning (day 1) (see also, Soeter and Kindt, 2011b). Note that the β-blocker (propranolol HCl) during memory retrieval (day 2) may also have suppressed a potential fear-enhancing effect of the yohimbine HCl manipulation (day 1). In any case, whereas the α_2-adrenergic drug delayed the process of extinction learning and triggered broader fear generalization 48 h later, the β-adrenergic receptor blocker (ie, propranolol HCl) during reconsolidation (day 2) selectively diminished the startle fear responding to the reactivated fear association along with its category-related information (day 3). Moreover, we observed a similar reduction in startle fear responding when the β-adrenergic receptor antagonist was administered after reactivation of the memory (ie, Experiment II), suggesting that the propranolol HCl manipulation before reactivation also affected the processes mediating reconsolidation (ie, Experiment I) (Nader et al, 2000). The present study employed stimuli of different 'valence' categories (fear-relevant vs fear-irrelevant) to verify whether the propranolol HCl manipulation was capable of neutralizing fear responding. That is, since fear-relevant stimuli are known to have an innate prepotency to elicit fear responses (Lovibond et al, 1994), we employed a fear-irrelevant cue as an additional control stimulus. The downside of this procedure is that the differential responding observed during acquisition (day 1) may simply be due to the emotional 'valence' of the stimuli employed (fear-relevant vs fear-irrelevant) rather than associative learning. Previously, we demonstrated in a fear conditioning paradigm that fear relevancy does not affect fear responding in the absence of associative learning (ie, first acquisition trial; Soeter and Kindt, 2011a). Again, we did not observe any difference in fear responding to the first trial of acquisition, indicating that the differential fear responding observed at the end of fear conditioning (day 1) was due to associative learning instead of the 'valence' of the cues. Thus, taken together, given that we observed a similar level of startle fear responding to the reactivated fear association and the fear-irrelevant control stimulus at retention testing (day 3), the β-adrenergic receptor blocker during reconsolidation apparently 'neutralized' the fear-arousing aspects of the 'associative' fear memory along with its category-related information (ie, generalization testing).

In line with our preceding studies, the α_2-adrenergic (Soeter and Kindt, 2011b) as well as β-adrenergic drug (Kindt et al, 2009; Soeter and Kindt, 2010, 2011b) did not affect the skin conductance responding and retrospective US expectancy ratings. Considering that online ratings direct the attention towards the CS–US relation (Baeyens et al, 1990; Lovibond and Shanks, 2002) and skin conductance responding is highly sensitive to attentional processes (Filion et al, 1991), the use of online ratings may previously have interfered with the measurement of electrodermal activity (Soeter and Kindt, 2010, 2011a, b). The current observation that the omission of online ratings does not differentially affect skin conductance responding corresponds to the view that electrodermal activity may primarily reflect the more cognitive level of contingency learning (ie, declarative knowledge) (Weike et al, 2007; but see, Schultz and Helmstetter, 2010). Studying human fear conditioning allows for the independent evaluation of both declarative knowledge and the fear response. In most fear conditioning studies, the conscious anticipation of an aversive stimulus (US) is associated with an increase in startle potentiation (Grillon and Davis, 1995; Lovibond and Shanks, 2002). However, there are also several observations showing that unawareness of a CS–US contingency does not preclude a startle fear response (eg, Weike et al, 2007). This indicates that the anticipation of an aversive stimulus is not a necessary condition to observe fear-potentiated startle responses. Our findings show that the anticipation of an US is also not a sufficient condition to generate fear-potentiated startle responses. As such, our results emphasize the concept of multiple memory systems and suggest a double dissociation between the emotional and cognitive representation of fear (Squire, 2005; LaBar and Cabeza, 2006; Weike et al, 2005, 2007). It should be noted that these findings do not imply that reconsolidation is restricted to the emotional expression of fear memory (ie, startle fear responding). In view of the hypothesized adaptive function of memory reconsolidation (Dudai, 2006, 2009; Lee, 2009), there is no a priori reason for assuming that some memory systems would not be sensitive to disrupting reconsolidation (Lee, 2009).

In our previous work (Kindt et al, 2009; Soeter and Kindt, 2010), the erasure of the fear response could also have resulted from a more diffuse effect of the propranolol HCl manipulation by reducing the fear-provoking aspects of the aversive consequence itself (US). In considering clinical implications, disrupting reconsolidation should not radically alter functional reactions to potentially dangerous situations (US), but selectively weaken the underlying maladaptive fear association (CS1–US). Our current observation that the non-reactivated stimulus (CS2) potentiated the startle fear responding at test (day 3) indicates that the participants still feared the US, but no longer the reactivated conditioned stimulus (CS1) (see also, Soeter and Kindt, 2011a). Apparently, the propranolol HCl manipulation selectively disrupted the CS1–US association at the emotional level given that the participants also expected the US when presented with the reactivated stimulus (CS1) at test (day 3). Taken together, these findings show that there is no causal link between the 'actual knowledge' of a fear association and its fear response, even though they often operate in parallel. Note that the fear-erasing effects were

not restricted to the feared cue itself (CS1), but instead generalized to category-related information not previously associated with the originally feared stimulus. The generalization of fear has been demonstrated to be dependent on the training-induced strength of the memory (Laxmi *et al*, 2003). Here, the strengthening of the fear memory trace by the α_2-adrenergic manipulation indeed triggered broader fear generalization (category-related word cues). Conversely, the β-adrenergic interference with reconsolidation left the memory trace too weak to yield a generalized fear response (Soeter and Kindt, 2011a). Several findings have implicated CREB phosphorylation in the formation (Josselyn *et al*, 2001; Davies *et al*, 2004) as well as generalization of fear memory (Han *et al*, 2008). Whereas yohimbine HCl is known to induce *p*CREB activation (Sun *et al*, 2010), the β-adrenergic receptor antagonist propranolol has been shown to inhibit noradrenaline-stimulated CREB phosphorylation (Thonberg *et al*, 2002). The present findings show the involvement of noradrenergic modulation in the generalization of human associative fear memory. Corroborating our previous study, the findings further demonstrate that the assimilation of individual memory items into a generalized schema may be dissociable for semantic and affective knowledge (Soeter and Kindt, 2011a). That is, upon exposure to the category-related information of the reactivated fear memory, the participants again predicted danger without a concomitant fear response.

While fear responses are very common in the aftermath of a traumatic event, the intensity of this initial fear responding is generally a poor indicator of symptom development or PTSD diagnosis (Rothbaum *et al*, 1992; Brewin *et al*, 2000; Murray *et al*, 2002). Rather, the impairment in the 'unlearning' of fear-related behavior as well as the generalization of fear to intrinsically safe stimuli (Blechert *et al*, 2007; Lissek and Grillon, 2010) has a strong relevance to the development of anxiety disorders such as PTSD. Hence, our data indicate that the noradrenaline level during or shortly after a traumatic experience may contribute to the etiology of post-traumatic stress disorder. At the same time, the present findings may have valuable therapeutic implications given that the strength of the fear memory did not act as a constraint on reconsolidation. It may be possible that the stimulation of the noradrenergic system during memory formation was not strong enough to prevent the disruption of reconsolidation (Suzuki *et al*, 2004; Wang *et al*, 2009). On the other hand, the yohimbine HCl manipulation yielded a resistance to fear extinction 48 h later and triggered broader fear generalization, indicating the former strengthening of the excitatory fear memory (Myers and Davis, 2002; Soeter and Kindt, 2011b). Moreover, preliminary evidence in trauma patients also revealed diminished trauma-relevant physiological responding following the β-adrenergic interference with reconsolidation (Brunet *et al*, 2008). The current observation that propranolol HCl can well be administered after reactivation of the fear memory (ie, Experiment II) further has implications in light of the demarcation between 'extinction' and 'reconsolidation' (Eisenberg *et al*, 2003; Pedreira and Maldonado, 2003; Suzuki *et al*, 2004). That is, when memory retrieval initiates extinction learning instead of reconsolidation, the propranolol HCl manipulation should be omitted to avoid interference with the consolidation of extinction training (Ouyang and Thomas, 2005; Mueller *et al*, 2008).

The 'process of extinction' and 'disrupting reconsolidation' are two approaches to diminish fear-related behavior. The extinction of fear not only leaves the original fear memory intact—thereby explaining the return of fear after apparently successful fear reduction (Bouton, 1993)—but may also be impaired by noradrenergic stimulation during the formation of the excitatory fear memory. Conversely, β-adrenergic receptor blockade during reconsolidation selectively 'deletes' the fear-arousing aspect of noradrenergic strengthened fear memory and—on top of that—undermines the generalization of fear responding. Given that fear generalization lies at the heart of many anxiety disorders, disrupting reconsolidation points to a promising strategy in reducing excessive fear responding.

DISCLOSURE

The authors declare no conflict of interest.

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Supplementary Information accompanies the paper on the Neuropsychopharmacology website (http://www.nature.com/npp)